Syndromic Surveillance Reporting (Optional)

07/08/2024 8:00 pm EDT

If you are engaged with a public health agency to submit syndromic surveillance data from an urgent care setting, you can attest to measure in your Healthmonix MIPSpro account.

Description

The MIPS-eligible clinician is in active engagement with a public health agency to submit syndromic surveillance data from an urgent care setting.

Definitions

Active Engagement - The MIPS-eligible clinician is in the process of moving towards sending "production data" to a public health agency or clinical data registry, or is sending production data to a public health agency (PHA) or clinical data registry (CDR).

Active engagement may be demonstrated in one of the following ways:

- **Option 1 – Pre-Production and Validation**: The MIPS-eligible clinician must first register to submit data with the PHA or, where applicable, the clinical data registry (CDR) to which the information is being submitted. Registration must be completed within 60 days after the start of the performance period, while awaiting an invitation from the PHA or CDR to begin testing and validation. MIPS-eligible clinicians that have registered in previous years do not need to submit an additional registration for subsequent performance periods. Upon completion of the initial registration, the MIPS-eligible clinician must begin the process of testing and validation of the electronic submission of data. The MIPS-eligible clinician must respond to requests from the PHA or, where applicable, the CDR within 30 days; failure to respond twice within a performance period would result in the MIPS-eligible clinician not meeting the measure.
- **Option 2 – Validated Data Production:** The MIPS-eligible clinician has completed testing and validation of the electronic submission and is electronically submitting production data to the PHA or CDR.

Production data – Refers to data generated through clinical processes involving patient care, and it is used to distinguish between data and “test data” which may be submitted for the purposes of enrolling in and testing electronic data transfers.

Reporting Requirements

The MIPS eligible clinician must attest YES to being in active engagement with a PHA to submit syndromic surveillance data from an urgent care setting.

Option 1/Option 2

In addition to submitting a response, MIPS-eligible clinicians must submit their level of active engagement, either Pre-production and Validation or Validated Data Production for each measure they report beginning with the performance period in CY 2023.

In your Healthmonix MIPSpro account, you can attest to the measure. Click **Save** when finished.

Syndromic Surveillance Reporting (PI_PHCDRR_2)

Complete:

- ○ This group qualifies under option 1 of active engagement public health agency to submit syndromic surveillance data from an urgent care setting.
- ○ This group qualifies under option 2 of active engagement with a public health agency to submit syndromic surveillance data from an urgent care setting.

Measure Details

Measure Title: Syndromic Surveillance Reporting

Measure ID: PI_PHCDRR_2

Objective: Public Health and Clinical Data Exchange

Description

The MIPS eligible clinician is in active engagement with a public health agency to submit syndromic surveillance data from an urgent care setting.

Definitions

Active engagement - The MIPS eligible clinician is in the process of moving towards sending "production data" to a public health agency or clinical data registry, or is sending production data to a public health agency (PHA) or clinical data registry (CDR).

Active engagement may be demonstrated in one of the following ways:

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>
> **Option 2 – Validated Data Production:** The MIPS eligible clinician has completed testing and validation of the electronic submission and is electronically submitting production data to the PHA or CDR.

Production data - Refers to data generated through clinical processes involving patient care, and it is used to distinguish between data and "test data" which may be submitted for the purposes of enrolling in and testing electronic data transfers.

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